

Tiffany Grider

List of Publications by Year in descending order

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Version: 2024-02-01

11
papers

307
citations

1040056

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1281871

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docs citations

11
times ranked

491
citing authors

#	ARTICLE	IF	CITATIONS
1	MicroRNAs as Biomarkers of Charcot-Marie-Tooth Disease Type 1A. <i>Neurology</i> , 2021, 97, e489-e500.	1.1	14
2	Transmembrane protease serine 5: a novel Schwann cell plasma marker for CMT1A. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 69-82.	3.7	25
3	A longitudinal study of CMT1A using Rasch analysis based CMT neuropathy and examination scores. <i>Neurology</i> , 2020, 94, e884-e896.	1.1	29
4	The audiologic profile of patients with Charcot-Marie Tooth neuropathy can be characterised by both cochlear and neural deficits. <i>International Journal of Audiology</i> , 2019, 58, 902-912.	1.7	12
5	Validation of MRC Centre MRI calf muscle fat fraction protocol as an outcome measure in CMT1A. <i>Neurology</i> , 2018, 91, e1125-e1129.	1.1	43
6	Coexistence of a T118M <i>PMP22</i> missense mutation and chromosome 17 (17p11.2-12) deletion. <i>Muscle and Nerve</i> , 2015, 52, 905-908.	2.2	4
7	Progressive Lower Extremity Weakness and Axonal Sensorimotor Polyneuropathy from a Mutation in <i>KIF5A</i> (c.611G>A;p.Arg204Gln). <i>Case Reports in Genetics</i> , 2015, 2015, 1-5.	0.2	4
8	Rare Manifestation of a c.290 C>T, p.Gly97Glu <i>VCP</i> Mutation. <i>Case Reports in Genetics</i> , 2015, 2015, 1-5.	0.2	16
9	Reduced neurofilament expression in cutaneous nerve fibers of patients with CMT2E. <i>Neurology</i> , 2015, 85, 228-234.	1.1	21
10	Genotype-phenotype characteristics and baseline natural history of heritable neuropathies caused by mutations in the <i>MPZ</i> gene. <i>Brain</i> , 2015, 138, 3180-3192.	7.6	80
11	Psychometrics evaluation of Charcot-Marie-Tooth Neuropathy Score (<i>CMTNSv2</i>) second version, using Rasch analysis. <i>Journal of the Peripheral Nervous System</i> , 2014, 19, 192-196.	3.1	59